

ADDITIVE FOR RUBBER COMPOSITION, ADDITIVE COMPOSITION
FOR RUBBER COMPOSITION, RUBBER COMPOSITION AND TIRE

BACKGROUND OF THE INVENTION

1. Field of the Invention

[0001] This invention relates to an additive for a rubber composition and an additive composition for a rubber composition capable of improving a processability
5 and productivity in the production of rubber articles without badly affecting properties of an uncured rubber and of a cured rubber as well as a rubber composition and a pneumatic tire using the same.

2. Description of Related Art

[0002] In the production of rubber articles such as tires, belts, hoses and the like,
10 kneading of a rubber composition is generally conducted in order for uniformly dispersing compounding ingredients and filler into starting rubber. And also, it is required to further conduct rubber milling for plasticizing the rubber composition several times in order to facilitate building operability at subsequent rubber processing step. However, such a repetition of the plasticizing operation is
15 unfavorable in view of the productivity of the rubber article. On the other hand, although it is effective to decrease the rubber milling number in order to improve the productivity of the rubber article, the uncured rubber composition is not sufficiently plasticized when the rubber milling number is merely decreased, which does not ultimately link to the improvement of the productivity.

[0003] To this end, it is possible to improve the processability of an uncured
20 rubber composition by adding a plasticizer, a processing aid or the like. Then, the building operability can be improved while decreasing the rubber milling number. In case of using a conventional plasticizer, a processing aid and the like, however, the degradation of properties of the uncured rubber composition or the cured rubber
25 composition is caused and it is actually difficult to decrease the rubber milling number. At the processing step in the production of the rubber article including natural rubber is particularly existent a polymer gel formed by entanglement of natural rubber molecular chains with each other, or by reaction of functional groups in isoprene chain of natural rubber with each other, or by reaction of such a functional
30 group with non-rubber ingredient in natural rubber, so that the rubber milling number increases and the degradation of the building operability is accompanied

therewith. And also, if the rubber milling number is increased for plasticizing an uncured rubber composition, the lowering of molecular weight in natural rubber is caused, which badly affects the properties of the cured rubber composition.

[0004] Therefore, it is demanded to improve the building operability without
5 degrading the properties of the uncured rubber composition or the cured rubber composition in the production of the rubber article particularly including natural rubber since early times.

[0005] In order to solve these problems, there are recently proposed some attempts. For instance, JP-A-11-209406 discloses that a quantity of polymer gel in
10 natural rubber is decreased by adding an aromatic polycarboxylic acid derivative to natural rubber. According to this method, however, it has been confirmed that blooming is apt to be easily caused in the cured rubber.

[0006] And also, JP-A-6-57040 and JP-A-4-20579 disclose that the process-
ability is improved by adding a specified ester. According to these methods,
15 however, peculiar odor is generated in the processing, which becomes a problem in view of the operating environment.

SUMMARY OF THE INVENTION

[0007] It is, therefore, an object of the invention to solve the aforementioned problems of the conventional techniques and to provide an additive for a rubber
20 composition and an additive composition for a rubber composition both being capable of improving the processability without degrading the properties of the rubber composition. It is another object of the invention to solve the problem in view of the operating environment and particularly decreasing the quantity of polymer gel in a rubber composition including natural rubber without lowering the
25 molecular weight of natural rubber to thereby improve the processability and the productivity and stabilize surface states of both uncured and cured rubber compositions as well as a rubber composition and a pneumatic tire using the same.

[0008] The inventors have discovered that slippage between rubber molecules is increased by using a partial ester between a specified aliphatic or aromatic polyvalent
30 carboxylic acid and a (poly)oxyalkylene derivative as an additive for a rubber composition without degrading the properties of a cured rubber composition and as a result the invention has been accomplished.

[0009] According to a first aspect of the invention, there is the provision of an

additive for a rubber composition consisting essentially of an ester selected from an ester of (i) an aliphatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group in its molecule, and an ester of (ii) an aromatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule.

[0010] According to a second aspect of the invention, there is the provision of an additive composition for a rubber composition comprising (a) a reinforcing filler and (b) an ester selected from an ester of (i) an aliphatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group in its molecule, and an ester of (ii) an aromatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule, in which a compounding ratio by weight of the component (a) to the component (b) is 70/30 to 30/70.

[0011] According to a third aspect of the invention, there is the provision of a rubber composition comprising a rubber ingredient and an ester selected from an ester of (i) an aliphatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group in its molecule, and an ester of (ii) an aromatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule.

[0012] According to a fourth aspect of the invention, there is the provision of a rubber composition comprising (A) a rubber ingredient and (B) an additive composition for a rubber composition consisting of (a) a reinforcing filler and (b) an ester selected from an ester of an aliphatic polyvalent carboxylic acid with a (poly)oxyalkylene derivative, and having at least one carboxyl group in its molecule, and an ester of an aromatic polyvalent carboxylic acid with a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule.

[0013] According to a fifth aspect of the invention, there is the provision of a pneumatic tire characterized by using a rubber composition defined in any one of the third and fourth aspects of the invention.

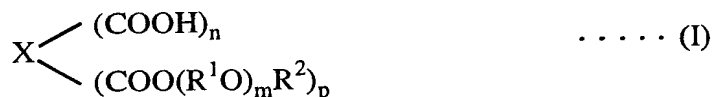
DETAILED DESCRIPTION OF THE INVENTION

[0014] The additive for a rubber composition according to the invention is an ester of an aliphatic polyvalent carboxylic acid with a (poly)oxyalkylene derivative,

and having at least one carboxyl group in its molecule, or an ester of an aromatic polyvalent carboxylic acid with a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule.

[0015] The former ester is preferable to be represented by the following formula

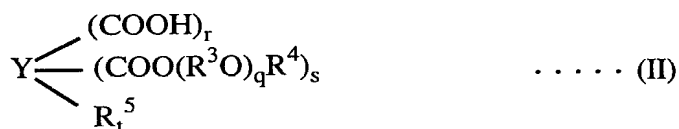
(I):



(wherein m is a number of not less than 1 indicating an average degree of polymerization, and each of n and p is an integer of not less than 1, and X is a saturated or unsaturated aliphatic chain, and R¹ is an alkylene group, and R² is a hydrocarbon group selected for the group consisting of an alkyl group, an alkenyl group, an alkylaryl group and an acyl group).

[0016] In the formula (I), it is favorable that X is an aliphatic chain having an unsaturated bond, and n = 1 and p = 1, and R¹ is an alkylene group having a carbon number of 2-4, and R² is an alkyl group or an alkenyl group having a carbon number of 2-28. More preferably, X is an aliphatic chain having an unsaturated bond and a carbon number of 2-8, and m is 1-10, and R¹ is an ethylene group or a propylene group, and R² is an alkyl group or an alkenyl group having a carbon number of 8-18.

[0017] The latter ester is preferable to be represented by the following formula (II):



(wherein q is a number of not less than 1 indicating an average degree of polymerization, and each of r and s is an integer of not less than 1, and t is 0 or an integer of not less than 1, and r+t+s is 6-8, and Y is an aromatic ring, and R³ is an alkylene group, R⁴ is a hydrocarbon group selected from the group consisting of an alkyl group, an alkenyl group, an alkylaryl group and an acyl group, and R⁵ is a hydrogen atom, an alkyl group or an alkenyl group).

[0018] In the formula (II), it is favorable that r+s is 2 or 3, and R³ is an alkylene group having a carbon number of 2-4, and R⁴ is an alkyl group or an alkenyl group having a carbon number of 2-28. More preferably, Y is a benzene ring, and r = 1

and $s = 1$, and R^3 is an ethylene group, and R^4 is an alkyl group or an alkenyl group having a carbon number of 2-28. Most preferably, q is 1-10, $r = 1$, $s = 1$, R^3 is an ethylene group and R^4 is an alkyl group or an alkenyl group having a carbon number of 8-18.

5 **[0019]** The additive for a rubber composition according to the invention is obtained by reacting (i) an aliphatic polyvalent carboxylic acid, in another word a bivalent or more aliphatic carboxylic acid or anhydride thereof, or (ii) an aromatic polyvalent carboxylic acid, in another word, a bivalent or more aromatic carboxylic acid or anhydride thereof with (iii) a (poly)oxyalkylene derivative.

10 **[0020]** As the polyvalent aliphatic carboxylic acid or the anhydride thereof (i), mention may be made of saturated aliphatic bivalent carboxylic acids such as succinic acid, succinic anhydride, glutaric acid, adipic acid, sebacic acid and the like; unsaturated aliphatic bivalent carboxylic acids such as maleic acid, maleic anhydride, fumaric acid, itaconic acid, itaconic anhydride, citraconic acid, citraconic
15 anhydride, alkenylsuccinic acid, alkenylsuccinic anhydride and the like; aliphatic trivalent carboxylic acid such as maleinized aliphatic acid or the like; and so on. As the component (i), bivalent carboxylic acid having an unsaturated bond or an anhydride thereof is favorable, and a maleic anhydride is most favorable. These polyvalent aliphatic carboxylic acids and anhydrides thereof may be used alone or in
20 admixture of tow or more.

[0021] As the polyvalent aromatic carboxylic acid (ii), mention may be made of bivalent aromatic carboxylic acids and their anhydrides such as phthalic acid, anhydrous phthalic acid, naphthalene dicarboxylic acid and the like; trivalent aromatic carboxylic acids and their anhydrides such as trimellitic acid, anhydrous
25 trimellitic acid and the like; tetravalent aromatic carboxylic acids and their anhydrides such as pyromellitic acid, anhydrous pyromellitic acid and the like; and so on. As the component (ii), bivalent or trivalent aromatic acids and anhydrides thereof are favorable in view of the cost, and anhydrous phthalic acid is particularly favorable. These polyvalent aromatic carboxylic acids and anhydrides thereof may
30 be used alone or in admixture of two or more.

[0022] As the (poly)oxyalkylene derivative (iii), mention may be made of compounds having a (poly)oxyalkylene group with an average polymerization degree of not less than 1, and having one or more hydroxyl groups, preferably

compounds having a (poly)oxyalkylene group with one or two hydroxyl groups, more particularly compounds having a (poly)oxyalkylene group and one hydroxyl group. As the component (iii), mention may be made of ether-type derivatives such as (poly)oxyalkylene alkyl ether and the like; ester-type derivatives such as (poly)oxyalkylene aliphatic acid monoester and the like; ether ester-type derivatives such as (poly)oxyalkylene glycerine aliphatic acid ester and the like; nitrogen-containing derivatives such as (poly)oxyalkylene aliphatic acid amide, (poly)oxyalkylene alkylamine and the like; and so on. Among them, the ether-type and ester-type derivatives are favorable, and the ether-type derivatives are particularly favorable.

[0023] As the ether-type (poly)oxyalkylene derivative, mention may be made of polyoxyalkylene aliphatic ethers such as polyoxyethylene lauryl ether, polyoxyethylene decyl ether, polyoxyethylene octyl ether, polyoxyethylene 2-ethylhexyl ether, polyoxyethylene polyoxypropylene lauryl ether, polyoxypropylene stearyl ether, polyoxyethylene oleyl ether and the like; polyoxyethylene aromatic ethers such as polyoxyethylene benzyl ether, polyoxyethylene alkylphenyl ether, polyoxyethylene benzylated phenyl ether and the like; and so on. Among them, the polyoxyalkylene aliphatic ether is favorable.

[0024] Further, polyoxyethylene alkyl or alkenyl ether is preferable. In this case, it is favorable that an average polymerization degree of polyoxyethylene is 1-10 and a carbon number of alkyl or alkenyl group is 8-18. Concretely, when polyoxyethylene is abbreviated as POE(n) wherein n is an average polymerization degree, there are mentioned POE(3) octyl ether, POE(4) 2-ethylhexyl ether, POE(3) decyl ether, POE(5) decyl ether, POE(3) lauryl ether, POE(8) lauryl ether, POE(1) stearyl ether and so on.

[0025] Moreover, the above (poly)oxyalkylene derivatives may be used alone or in admixture of two or more.

[0026] In the ester of the component (i) and the component (iii) according to the invention, the component (i) as a starting material may be included in an amount of not more than 10% by weight, preferably not more than 5% by weight, or the component (iii) as a starting material may be included in an amount of not more than 40% by weight, preferably not more than 20% by weight. As such an ester, mention may be made of mono(POE(3) lauryl ether) maleic acid ester, mono(POE(3) 2-ethylhexyl

ether) maleic acid ester and so on.

[0027] The additive composition for a rubber composition according to the invention comprises (a) a reinforcing filler and (b) an ester selected from an ester of (i) an aliphatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group in its molecule, and an ester of (ii) an aromatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule, in which a compounding ratio by weight of the component (a) to the component (b) is preferable to be 70/30 to 30/70 in view of the operability. When the compounding ratio is less than 30/70, the handling may become difficult, while when it exceeds 70/30, the improvement of the operability may not be satisfactorily obtained. In this point, the compounding ratio by weight is more preferable to be within a range of 60/40 to 40/60.

[0028] The reinforcing filler as the component (a) is not particularly restricted, but is favorable to be carbon black, inorganic fillers, such as silica, aluminum hydroxide, talc, clay and the like. Among them, carbon black, silica and aluminum hydroxide are preferable. Especially, as a carbon black, HAF, ISAF and SAF are preferable, while silica is preferable to have an N_2SA (nitrogen adsorption specific surface area) of 50-250 m^2/g and a DBP (dibutyl phthalate absorption) of 150-300 ml/100 g.

[0029] Further, the additive composition for a rubber composition according to the invention may contain at least one compound selected from (poly)oxyalkylene derivatives, alcohols and aliphatic esters as a component (c). In this case, a compounding amount of the component (c) is preferable to be not more than 10 parts by weight per 100 parts by weight of total amount of the components (a) and (b). When the compounding amount exceeds 10 parts by weight, merits by the addition of a large amount of the component (c) are not substantially obtained. These compounds may be used alone or in admixture of two or more as the component (c).

[0030] The component (c) is favorable to have a melting point of not higher than 20°C and a boiling point of not lower than 150°C. When the melting point exceeds 20°C, it is difficult to control powdery dust in the production of the additive composition or a rubber composition using the same, while when the boiling point is lower than 150°C, the safeness in the rubber processing can not be sufficiently ensured.

[0031] The (poly)oxyalkylene derivative as the component (c) includes derivatives of ether-type, ester-type, ether ester-type, nitrogen containing-type and so on likewise the above component (iii). Among them, ether-type or ester-type polyoxyalkylene derivative is favorable. Particularly, polyoxyethylene derivatives are preferable. Concretely, when polyoxyethylene is abbreviated as POE(n) and n is an average polymerization degree, there are mentioned POE(3) lauryl ether, POE(5) decyl ether, POE(4) oleyl ether, POE(10) monolaurate, POE(10) monooleate, ethylene glycol dioleate, POE(9) dicaprylate and so on.

[0032] The alcohol as the component (c) is not particularly restricted, but it is preferable to be an aliphatic alcohol having a carbon number of 8-20. For example, there are mentioned octyl alcohol, 2-ethylhexyl alcohol, decyl alcohol, lauryl alcohol, oleyl alcohol, stearyl alcohol, ethylene glycol, capryl alcohol and so on. And also, esters of these alcohols are preferably used as the aliphatic ester as the component (c).

[0033] The additive composition for a rubber composition according to the invention is obtained by merely mixing the above components in a mixer. As the mixer, mention may be made of a Henschel mixer, a Nauta mixer (trade name, made by Hosokawa Micron Co., Ltd.), a tumbler mixer and so on. The mixing may be carried out by selecting revolution number, agitating temperature, agitating time and the like within optimum ranges in accordance with the kind of the mixer used and the ingredient to be mixed.

[0034] The first rubber composition according to the invention is obtained by compounding a rubber ingredient with the additive for a rubber composition as previously mentioned, i.e. the ester of (i) an aliphatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group in its molecule, or the ester of (ii) an aromatic polyvalent carboxylic acid with (iii) a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule. If necessary, a reinforcing filler similar to those previously described may be compounded. As the reinforcing filler, carbon black, silica and aluminum hydroxide are favorable.

[0035] In this case, it is favorable that the compounding amount of the reinforcing filler is 10-85 parts by weight per 100 parts by weight of the rubber ingredient. And also, it is favorable that the compounding amount of the ester is

0.1-10 parts by weight, preferably 0.5-5 parts by weight per 100 parts by weight of the rubber ingredient. When the amount of the ester is not less than 0.1 part by weight, the processability is improved, while when it is not more than 10 parts by weight, the rubber properties are held and the cost becomes desirable.

5 **[0036]** The second rubber composition according to the invention is obtained by compounding (A) a rubber ingredient with (B) the additive composition for a rubber composition as previously mentioned, i.e. the additive composition consisting of (a) the reinforcing filler and (b) the ester of an aliphatic polyvalent carboxylic acid with a (poly)oxyalkylene derivative, and having at least one carboxyl group in its
10 molecule, or the ester of an aromatic polyvalent carboxylic acid with a (poly)oxyalkylene derivative, and having at least one carboxyl group bonded to an aromatic ring in its molecule. If necessary, at least one of a (poly)oxyalkylene derivative, an alcohol and an aliphatic acid ester thereof may be compounded.

[0037] In the component (B) used in the above second rubber composition, the
15 compounding ratio by weight of the component (a) to the component (b) is 70/30-30/70, preferably 60/40-40/60. And also, the compounding amount of the component (B) in the second rubber composition is 0.1-10 parts by weight, preferably 0.5-5 parts by weight per 100 parts by weight of the rubber ingredient.

[0038] In any case, the rubber composition is obtained, for example, by
20 kneading the above components to be compounded.

[0039] When the rubber ingredient includes natural rubber, the rubber compositions according to the invention can reduce a polymer gel quantity of natural rubber without decreasing the molecular weight of natural rubber and increases the slippage between rubber molecules to improve the building operability
25 and control the degradation of the properties in the uncured and cured rubber compositions. On the other hand, when the rubber ingredient does not include natural rubber, the slippage between rubber molecules is increased to improve the building operability and also the decrease in molecular weight of rubber component is not caused and hence the degradation of the properties in the uncured and cured
30 rubber compositions is not caused. In any case, according to the invention, the excellent rubber processability can be obtained without badly affecting the properties of either uncured or cured rubber composition.

[0040] The rubber ingredient used in the rubber compositions according to the

invention is not particularly restricted, but is favorable to be natural rubber alone or a blend of natural rubber and synthetic rubber(s). As the synthetic rubber, mention may be made of styrene-butadiene rubber (SBR), butadiene rubber (BR), isoprene rubber (IR), butyl rubber (IIR), halogenated butyl rubber (X-IIR), ethylene-propylene-diene terpolymer rubber (EPDM) and a mixture thereof.

[0041] As the reinforcing filler used in the rubber composition, use may be made of carbon blacks of HAF, ISAF, SAF and the like, inorganic fillers such as silica, aluminum hydroxide, clay talc and so on as previously mentioned. They may be used alone or in admixture of two or more. Moreover, when an inorganic filler is compounded in the rubber composition, the dispersion improving effect through the inorganic filler is obtained, and such an effect becomes remarkable as the silica content increases.

[0042] Furthermore, the rubber composition according to the invention may properly be compounded with the other additives usually used in the rubber field such as sulfur, vulcanization accelerator, processing oil, antioxidant and so on, if necessary.

[0043] The rubber composition according to the invention is obtained by kneading the aforementioned components in a kneading machine such as rolls, internal mixer or the like and subjected to a given building process, which can be applied to not only application of tire member such as tread, under tread, carcass, sidewall, bead portion or the like but also application of the other industrial goods such as rubber vibration isolator, belt, hose and the like. Preferably, the above rubber composition is used as a rubber for tire tread.

[0044] The pneumatic tire according to the invention is manufactured by using the above rubber composition according to usual manner. That is, the rubber composition according to the invention containing, if necessary, various chemicals is extruded into various members for the tire at an uncured stage and shaped into a green tire on a tire building machine according to the usual manner. This green tire is heated under pressure in a vulcanizing machine to form a given tire. In the thus obtained tire, the rubber properties are not degraded, and also the productivity is excellent because the processability of the rubber composition is good. Moreover, the tire according to the invention can be used by filling a gas such as air, nitrogen or the like in the inside of the tire.

[0045] The following examples are given in illustration of the invention and are not intended as limitations thereof.

[0046] Various measurements in the examples are carried out by the following methods.

5 **[0047]** Mooney viscosity (ML_{1+4})

As an evaluation of processability of a rubber composition, an uncured rubber sample compounded and kneaded with a vulcanizing system is preheated at 130°C for 1 minute and a rotor is rotated for 4 minutes to measure a Mooney viscosity by using a Mooney viscometer SMV201 made by Shimadzu Seisakusho.

10 The measured value is represented by an index on the basis that Comparative Example 1, 2 or 3 is 100. The larger the index value, the better the processability.

[0048] Mooney scotch time

As an evaluation of processing stability of a rubber composition, an uncured rubber sample compounded and kneaded with a vulcanizing system is
15 preheated at 130°C for 1 minute by using the same apparatus as used in the measurement of the Mooney viscosity and then a time taken for a rise in Mooney viscosity by 5 units from a minimum value V_m at the start of preheating is measured according to JIS K6300-1994. The measured value is represented by an index on the basis that Comparative Example 3 is 100. The smaller the index value, the
20 better the processing stability.

[0049] Vulcanizing property (T90)

As an evaluation of adequate vulcanizing property of a rubber composition, a value of 90% vulcanizing time (T90) is measured at 150°C according to vibration type vulcanization test of JIS K6300-1994. The measured value is
25 represented by an index on the basis that Comparative Example 3 is 100. The smaller the index value, the shorter the vulcanizing time and the better the vulcanization productivity.

[0050] Odor

An odor felt in weighing and in kneading is evaluated by 10 workers
30 according to the following standard:

○: all of 10 workers do not feel odor

△: 1-9 workers feel some odor

X: all of 10 workers feel odor

[0051] Tensile strength at break, Elongation at break

The tensile strength at break and elongation at break are measured by using a dumbbell No. 3 specimen according to a test method of JIS K6251-1993.

The measured values are represented by an index on the basis that Comparative

- 5 Example 1, 2, 3 or 11 is 100. The larger the index value, the larger the tensile strength at break and the elongation at break.

[0052] Rebound resilience

A given weight is freely dropped from a given height onto a surface of cured sample to measure a jumping height of the weight (percentage to the original
10 height) according to a test method of JIS K6301-1995. The measured value is represented by an index on the basis that Comparative Example 1, 2, 3 or 11 is 100. The larger the index value, the larger the rebound resilience.

[0053] Wear resistance

- 15 A worn quantity is measured by means of a Lambourn abrasion tester according to JIS K6264-1993. The measured value is represented by an index on the basis that Comparative Example 1, 2, 3 or 11 is 100. The larger the index value, the better the wear resistance.

[0054] Presence or absence of blooming

- 20 After a cured rubber specimen after the vulcanization is left to stand in an atmosphere of 5°C for 12 weeks, toluene-soluble matter on the surface of the specimen is analyzed by means of a gas chromatograph and a mass spectrometer to measure presence or absence of a substance precipitated from an additive composition for a rubber composition or resulted from such an additive composition. Moreover, the measuring conditions in the gas chromatograph are that an injection
25 temperature is 280°C, and a column is held at 70°C for 1 minute and heated up to 300°C at a rate of 25°C/min and held at 300°C for 10 minutes, and helium is used as a carrier gas and a flow rate of the carrier gas is 1 ml/min, while the measuring condition in the mass spectrometer is that a scanning range is M/Z:25-500. As a result, the presence or absence of blooming is evaluated as follows.

- 30 ○: the precipitation of the substance is not confirmed

X: the precipitation of the substance is confirmed

[0055] Examples 1-7, Comparative Example 1

In a Banbury mixer are kneaded 100 parts by weight of natural rubber

(RSS#3), an ester as a component (b) of an amount per 100 parts by weight of rubber ingredient shown in Table 1 and a kind shown in Table 2, 55 parts by weight of carbon black HAF, 2 parts by weight of stearic acid, 1 part by weight of an antioxidant, 6PPD (N-phenyl-N'-(1,3-dimethylbutyl)-p-phenylene diamine) and 1 part by weight of a wax at a starting temperature of 70°C and a revolution number of 70 rpm for 4 minutes. After the resulting mixture is sufficiently cooled in air at room temperature, it is further kneaded with 3 parts by weight of zinc white, 1 part by weight of a vulcanization accelerator, DM (dibenzothiazyl disulfide), 1 part by weight of a vulcanization accelerator, CBS (N'-dicyclohexyl-2-benzothiazyl sulfenamide) and 2 parts by weight of sulfur at a starting temperature of 70°C and a revolution number of 70 rpm in a Banbury mixer for 1 minute. Moreover, an ester as the component (b) is not used in Comparative Example 1.

[0056] The Mooney viscosity is measured with respect to the resulting uncured rubber compositions. And also, such an uncured rubber composition is cured at 150°C for 30 minutes and then the tensile strength at break, elongation at break, rebound resilience and wear resistance are measured with respect to the resulting cured rubber to obtain results as shown in Table 1. Comparative Example 1 is the control in all the properties for all of Examples 1 to 7.

[0057]

Table 1

	Example 1	Example 2	Example 3	Example 4	Example 5	Example 6	Example 7	Comparative Example 1
Kind of component (b)	A	A	A	A	A	B	C	none
Amount of component (b) (phr)	0.05	0.3	3	8	15	3	3	—
Mooney viscosity	106	110	115	118	121	114	113	100
Tensile strength at break	101	99	100	98	98	101	99	100
Elongation at break	102	102	98	99	100	98	98	100
Rebound resilience	98	99	98	98	97	99	99	100
Wear resistance	98	97	99	100	98	97	99	100

[0058] As the ester of the component (b) are used compounds A, B and C having the following formula (III):



wherein m, R¹ and R² are shown in Table 2.

[0059]

Table 2

Compound	m	R ¹	R ²	Name
A	3	ethylene group	lauryl group	mono(POE(3) laurylether) maleic acid ester
B	3	propylene group	lauryl group	mono(POP(3) laurylether) maleic acid ester
C	3	ethylene group	2-ethylhexyl group	mono(POE(3)2-ethylhexyl ether) maleic acid ester

[0060] Examples 8-14, Comparative Example 2

An additive composition (B) for a rubber composition is previously prepared by mixing a component (b) of kind and amount shown in Table 3 with silica (Nipsil VN3, trade name, made by Nippon Silica Kogyo Co., Ltd.) at a mixing ratio of component (b):silica = 6:4.

[0061] In a Banbury mixer are kneaded 100 parts by weight of styrene-butadiene copolymer rubber (137.5 parts by weight of SBR1712, trade name, made by JSR Corporation, an oil-extended rubber containing 37.5 phr of aromatic oil), the above prepared additive composition, 30 parts by weight of carbon black ISAF, 30 parts by weight in total of silica (the same as silica used in the preparation of the additive composition), 1 part by weight of stearic acid, 1 part by weight of an antioxidant, 6PPD (N-phenyl-N'-(1,3-dimethylbutyl)-p-phenylene diamine), 1 part by weight of a wax and 3 parts by weight of a silane coupling agent (Si69, trade name, made by Degussa AG) at a starting temperature of 70°C and a revolution number of 70 rpm for 4 minutes.

[0062] After the resulting mixture is sufficiently cooled in air at room temperature, it is further kneaded with 3 parts by weight of zinc white, 1 part by weight of a vulcanization accelerator, DM (dibenzothiazyl disulfide), 1 part by weight of a vulcanization accelerator, D (diphenyl guanidine), 1 part by weight of a vulcanization accelerator, CBS (N-cyclohexyl-2-benzothiazyl sulfenamide) and 2 parts by weight of sulfur at a starting temperature of 70°C and a revolution number of

70 rpm in a Banbury mixer for 1 minute. Moreover, an ester as the component (b) is not used in Comparative Example 2.

- [0063]** The Mooney viscosity is measured with respect to the resulting uncured rubber composition. And also, such an uncured rubber composition is cured at 150°C for 30 minutes and then the strength at break, elongation at break, rebound resilience and wear resistance are measured with respect to the resulting cured rubber to obtain results as shown in Table 3. Comparative Example 2 is the control in all the properties for all of Examples 8 to 14.

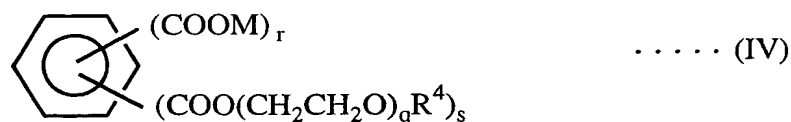
[0064]

Table 3

	Example 8	Example 9	Example 10	Example 11	Example 12	Example 13	Example 14	Comparative Example 2
Kind of component (b)	A	A	A	A	A	B	C	none
Amount of component (b) (phr)	0.05	0.3	3	8	15	3	3	—
Mooney viscosity	104	108	114	116	119	113	113	100
Tensile strength at break	101	100	99	98	99	100	99	100
Elongation at break	102	103	97	99	102	96	99	100
Rebound resilience	98	100	97	98	98	98	98	100
Wear resistance	98	98	98	100	100	95	99	100

- [0065]** As seen from the results of Tables 1 and 3, all of the rubber compositions according to the invention are considerably excellent in the processability of uncured rubber and maintain good properties of cured rubber without substantially degrading them.

[0066] In the following examples and comparative example, a compound represented by the following formula (IV):



- (wherein q, r, s, M, and R⁴ are shown in Table 4) is used as an additive for a rubber composition.

[0067]

Table 4

Carboxylic acid derivative			q	r	s	Kind of M	Kind of R ⁴
Additive for rubber composition	For Examples	I	3	1	1	H	lauryl
		II	3	1	1	H	decyl
		III	4	1	1	H	2-ethylhexyl
		IV	4	1	1	H	oleyl
		V	6	1	1	H	stearyl
		VI	6	2	1	H	stearyl
		VII	1	2	1	H	2-ethylhexyl
		VIII	12	2	1	H	octyl
	For Comparative Examples	IX	3	0	2	—	lauryl
		X	0	1	1	H	stearyl
		XI	0	1	1	H	decyl
		XII	0	1	1	H	oleyl
		XIII	3	1	1	Zn	lauryl

[0068] Examples 15-27, Comparative Examples 3-9

An uncured rubber composition is prepared by compounding 100 parts by weight of rubber ingredient shown in Table 5 with the kind and amount of the above additive for a rubber composition shown in Table 5, 50 parts by weight of carbon black ISAF, 2 parts by weight of stearic acid, 3 parts by weight of zinc white, 1 parts by weight of an antioxidant 6 PPD, 1.5 parts by weight of sulfur and 1 part by weight of a vulcanization accelerator CBS. The resulting uncured rubber composition is kneaded 3 times. Moreover, the kneading number in Comparative Example 3 is 4 times.

[0069] With respect to the thus obtained uncured rubber composition, the processability (Mooney viscosity), processing stability (Mooney scorch time), vulcanizing property (T90) and odor are evaluated to obtain results as shown in Table 5. Comparative Example 3 is set to be control in the properties for all of the examples and the comparative examples.

[0070]

Table 5

Uncured rubber	Rubber ingredient (A)		Additive for rubber composition		Number of kneading steps	Mooney viscosity	Mooney scorch time	Vulcanizing property T90	Odor
	natural rubber	SBR	Kind	parts by weight					
	parts by weight	parts by weight							
Example 15	100	0	I	1	3	102	92	95	○
Example 16	100	0	I	2	3	92	91	95	○
Example 17	100	0	I	5	3	88	103	91	○
Example 18	100	0	II	2	3	95	92	95	○
Example 19	100	0	III	2	3	95	90	91	○
Example 20	100	0	IV	2	3	99	99	98	○
Example 21	100	0	V	2	3	101	100	98	○
Example 22	100	0	VI	2	3	102	97	102	○
Example 23	100	0	VII	2	3	100	96	95	○
Example 24	100	0	VIII	2	3	99	95	98	○
Example 25	50	50	I	2	3	99	99	98	○
Example 26	50	50	II	2	3	100	98	97	○
Example 27	100	0	I	0.5	3	105	97	98	○
Comparative Example 3	100	0	—	0	4	100	100	100	○
Comparative Example 4	100	0	—	0	3	108	97	97	○
Comparative Example 5	100	0	IX	2	3	109	96	99	○
Comparative Example 6	100	0	X	2	3	91	90	93	○
Comparative Example 7	100	0	XI	2	3	94	92	93	×
Comparative Example 8	100	0	XII	2	3	104	95	93	×
Comparative Example 9	100	0	XIII	2	3	110	97	96	○

[0071] The rubber composition of Comparative Example 4 is poor in the processability as compared with the rubber composition of Comparative Example 3 having more of kneading number. On the contrary, when Examples 15-27 are compared with Comparative Example 3, the processability is improved even if the kneading number is decreased and the properties of the uncured rubber composition are maintained.

[0072] As seen from Examples 25 and 26, the invention is effective to the blend of natural rubber and synthetic rubber in addition to natural rubber alone.

10 [0073] In Comparative Examples 6-8, the kind of the additive for a rubber

composition is an alkyl ester, but is not (poly)oxyalkylene ester as defined in the invention, so that the working environment is degraded due to the odor although the degradation of the properties of the uncured rubber are not caused as shown in Comparative Examples 7 and 8 or the blooming of cured rubber becomes large as shown in the following Comparative Examples 12 and 13. Further, Comparative Examples 5 and 9 in which all of carboxyl groups are changed into ester and/or metal salt do not exhibit the effect on the properties such as processability and the like, from which it is clear that carboxyl group is essential.

[0074] Production Examples 1-11 (Additive composition for a rubber composition)

10 An additive composition for a rubber composition is prepared by mixing silica, carbon black as a component (a) with an ester of an aromatic polyvalent carboxylic acid and a (poly)oxyalkylene derivative as a component (b) and a (poly)oxyalkylene derivative or a polyvalent alcohol aliphatic acid ester having a melting point of not higher than 20°C and a boiling point of not lower than 150°C as a component (c) according to a compounding recipe as shown in Table 6 in a Henschel mixer while stirring. The property of the resulting additive composition is also shown in Table 6.

[0075]

Table 6

Production Example	Component (a)		Component (b)		Component (c)			Property
	kind	parts by weight	kind	parts by weight	kind	melting point (°C)	parts by weight	
1	silica	30	I	70	—	—	0	powder
2	silica	40	II	60	—	—	0	powder
3	silica	35	I	65	ethylene glycol oleate	-6	2	granulated powder
4	silica	45	I	55	ethylene glycol oleate	-6	5	granulated powder
5	silica	60	I	40	ethylene glycol oleate	-6	10	granulated powder
6	silica	45	I	55	POE(3)lauryl ether	-2	5	granulated powder
7	silica	45	I	55	POE(10)dicaprylate	0	5	granulated powder
8	silica	45	I	55	POE(10)monooleate	0	5	granulated powder
9	carbon black	65	I	35	—	—	0	powder
10	silica	65	II	35	ethylene glycol oleate	-6	5	granulated powder
11	silica	25	I	75	—	—	0	clay-like

[0076] The additive compositions of Production Examples 1-10 according to the invention are solid and are easy to handle. Particularly, in the additive compositions of Production Examples 3-8, 10, since the component (c) is added, the powdery duct is hardly scattered because these compositions are rendered into the granulated powder being more advantageous in the handling. On the contrary, the additive composition of Production Example 11 is clay-like and difficult in the handling.

[0077] Examples 28-41

100 parts by weight of rubber ingredient shown in Table 7 is kneaded with the additive compositions of Production Examples 1-10 of kind and amount shown in Table 7 (compounding amount is shown as an amount of component (b) in the composition), 50 parts by weight of carbon black ISAE, 2 parts by weight of stearic acid, 3 parts by weight of zinc white and 1 part by weight of an antioxidant 6 PPD, and then 1.5 parts by weight of sulfur and 1 part by weight of a vulcanization accelerator CBS as a vulcanizing additive are added and then the kneading is carried out at the number shown in Table 7 to obtain a rubber composition.

[0078] With respect to the thus obtained rubber compositions, the Mooney viscosity (processability), Mooney scorch time, vulcanizing property and odor are evaluated to obtain results as shown in Table 7.

[0079] Example 42

The same procedure as in Example 31 is repeated except that 25 parts by weight of carbon black and 25 parts by weight of silica are used instead of 50 parts by weight of carbon black in Example 31 and the same evaluations as in Example 31 are carried out to obtain results as shown in Table 7.

[0080]

Table 7

Uncured rubber	Rubber ingredient (A)		Additive composition for rubber (B)		Kneading number	Mooney viscosity	Mooney scorch time	Vulcanizing property T90	Odor
	natural rubber	SBR	Kind	component (b)					
	parts by weight	parts by weight		PHR*					
Example 28	100	0	Production Example 1	2	3	92	98	97	○
Example 29	100	0	Production Example 2	2	3	91	98	98	○
Example 30	100	0	Production Example 3	2	3	92	97	99	○
Example 31	100	0	Production Example 4	2	3	92	96	97	○
Example 32	100	0	Production Example 4	5	3	87	94	93	○
Example 33	100	0	Production Example 5	2	3	94	98	99	○
Example 34	100	0	Production Example 6	2	3	92	95	96	○
Example 35	100	0	Production Example 7	2	3	92	94	94	○
Example 36	100	0	Production Example 8	2	3	93	95	95	○
Example 37	100	0	Production Example 9	2	3	95	98	99	○
Example 38	100	0	Production Example 10	2	3	91	93	95	○
Example 39	50	50	Production Example 4	2	3	93	96	98	○
Example 40	50	50	Production Example 10	2	3	93	95	99	○
Example 41	100	0	Production Example 4	0.5	3	98	100	99	○
Example 42	100	0	Production Example 4	2	3	88	101	102	○

Note) *:compounding amount by weight per 100 parts by weight of rubber ingredient

- [0081] As seen from the results in Table 7, the same results as in Examples 15-27 can be said to be obtained even in Examples 28-41 using the previously prepared additive composition for rubber (B) containing the component (b). And also, according to the invention, the reinforcing filler is not restricted to carbon black as seen from Example 42.

[0082] Examples 43-48, Comparative Examples 10-13

The uncured rubber compositions shown in Table 8 are selected from the rubber compositions shown in Tables 5 and 7, and cured at 145°C for 33 minutes to obtain cured rubber samples. With respect to the thus obtained cured rubber samples, the tensile strength at break, elongation at break, rebound resilience, wear resistance and blooming are measured to obtain results as shown in Table 8. Comparative Example 11 is set to be control.

[0083]

Table 8

Cured rubber composition	No. of uncured rubber composition	Tensile strength at break	Elongation at break	Rebound resilience	Wear resistance	Blooming
Example 43	Example 16	101	100	98	100	○
Example 44	Example 18	100	102	98	100	○
Example 45	Example 25	100	100	99	100	○
Example 46	Example 26	100	100	99	100	○
Example 47	Example 31	101	100	99	100	○
Example 48	Example 38	101	100	99	100	○
Comparative Example 10	Comparative Example 3	98	98	97	97	○
Comparative Example 11	Comparative Example 4	100	100	100	100	○
Comparative Example 12	Comparative Example 6	98	99	95	99	×
Comparative Example 13	Comparative Example 8	100	100	98	100	×

[0084] As seen from Examples 43-48, even if the kneading number is decreased, the properties of the cured rubber are not badly affected and the blooming problem is not caused. However, the blooming is caused on the rubber surface in Comparative Examples 12 and 13 using no ester as defined in the invention.

[0085] As mentioned above, when the specified ester according to the invention is compounded with rubber component, even if the kneading number is decreased, the rubber properties are not substantially degraded over a period from all kneading steps to the production of rubber article.

[0086] By using the additive for a rubber composition and the additive composition for a rubber composition according to the invention, the processability of an uncured rubber composition is improved without degrading the properties of a

5 pneumatic tires.

Year	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030	2031	2032	2033	2034	2035	2036	2037	2038	2039	2040	2041	2042	2043	2044	2045	2046	2047	2048	2049	2050	2051	2052	2053	2054	2055	2056	2057	2058	2059	2060	2061	2062	2063	2064	2065	2066	2067	2068	2069	2070	2071	2072	2073	2074	2075	2076	2077	2078	2079	2080	2081	2082	2083	2084	2085	2086	2087	2088	2089	2090	2091	2092	2093	2094	2095	2096	2097	2098	2099	2100
1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030	2031	2032	2033	2034	2035	2036	2037	2038	2039	2040	2041	2042	2043	2044	2045	2046	2047	2048	2049	2050	2051	2052	2053	2054	2055	2056	2057	2058	2059	2060	2061	2062	2063	2064	2065	2066	2067	2068	2069	2070	2071	2072	2073	2074	2075	2076	2077	2078	2079	2080	2081	2082	2083	2084	2085	2086	2087	2088	2089	2090	2091	2092	2093	2094	2095	2096	2097	2098	2099	2100	